

BRG1 Rabbit mAb

Catalog # AP75169

Product Information

Application WB, IHC-P, IHC-F, IP, ICC

Primary Accession Q3TKT4

Reactivity Human, Mouse, Rat

Host Rabbi

Clonality Monoclonal Antibody

Calculated MW 181427

Additional Information

Gene ID 20586

Other Names Smarca4

Dilution WB~~1/500-1/1000 IHC-P~~N/A IHC-F~~N/A IP~~N/A ICC~~N/A

Format Liquid

Protein Information

Name Smarca4

Synonyms Baf190a, Brg1, Snf2b, Snf2l4

Function ATPase involved in transcriptional activation and repression of select genes

by chromatin remodeling (alteration of DNA-nucleosome topology).

Component of SWI/SNF chromatin remodeling complexes that carry out key enzymatic activities, changing chromatin structure by altering DNA-histone

contacts within a nucleosome in an ATP-dependent manner

(PubMed: <u>17640523</u>). Component of the CREST-BRG1 complex, a multiprotein

complex that regulates promoter activation by orchestrating the

calcium-dependent release of a repressor complex and the recruitment of an activator complex. In resting neurons, transcription of the c-FOS promoter is inhibited by SMARCA4-dependent recruitment of a phospho-RB1-HDAC repressor complex. Upon calcium influx, RB1 is dephosphorylated by calcineurin, which leads to release of the repressor complex. At the same time, there is increased recruitment of CREBBP to the promoter by a CREST-dependent mechanism, which leads to transcriptional activation. The

CREST-BRG1 complex also binds to the NR2B promoter, and

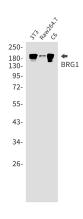
activity-dependent induction of NR2B expression involves the release of HDAC1 and recruitment of CREBBP (By similarity). Belongs to the neural progenitors-specific chromatin remodeling complex (npBAF complex) and the neuron-specific chromatin remodeling complex (nBAF complex). During neural development, a switch from a stem/progenitor to a postmitotic

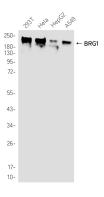
chromatin remodeling mechanism occurs as neurons exit the cell cycle and become committed to their adult state. The transition from proliferating neural stem/progenitor cells to postmitotic neurons requires a switch in subunit composition of the npBAF and nBAF complexes. As neural progenitors exit mitosis and differentiate into neurons, npBAF complexes which contain ACTL6A/BAF53A and PHF10/BAF45A, are exchanged for homologous alternative ACTL6B/BAF53B and DPF1/BAF45B or DPF3/BAF45C subunits in neuron-specific complexes (nBAF). The npBAF complex is essential for the self- renewal/proliferative capacity of the multipotent neural stem cells. The nBAF complex along with CREST plays a role in regulating the activity of genes essential for dendrite growth. SMARCA4/BAF190A may promote neural stem cell self-renewal/proliferation by enhancing Notchdependent proliferative signals, while concurrently making the neural stem cell insensitive to SHH-dependent differentiating cues (PubMed: 17640523). Acts as a corepressor of ZEB1 to regulate E-cadherin transcription and is required for induction of epithelial-mesenchymal transition (EMT) by ZEB1 (By similarity). Binds via DLX1 to enhancers located in the intergenic region between DLX5 and DLX6 and this binding is stabilized by the long non-coding RNA (IncRNA) Evf2 (PubMed:26138476). Binds to RNA in a promiscuous manner (PubMed: <u>26138476</u>). In brown adipose tissue, involved in the regulation of thermogenic genes expression (PubMed:34910916).

Cellular Location

Nucleus {ECO:0000255 | PROSITE-ProRule:PRU00549, ECO:0000269 | PubMed:26138476}. Note=Colocalizes with long non-coding RNA Evf2 in nuclear RNA clouds (PubMed:26138476). Localizes to sites of DNA damage (By similarity). {ECO:0000250 | UniProtKB:P51532, ECO:0000269 | PubMed:26138476}

Images





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